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NEWS	13	AUG 02 STN User Update to be held August 22 in conjunction with the 228th ACS National Meeting
NEWS	14	AUG 02 The Analysis Edition of STN Express with Discover! (Version 7.01 for Windows) now available
NEWS	15	AUG 04 Pricing for the Save Answers for SciFinder Wizard within STN Express with Discover! will change September 1, 2004
NEWS EXPRESS		JULY 30 CURRENT WINDOWS VERSION IS V7.01, CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP), AND CURRENT DISCOVER FILE IS DATED 26 APRIL 2004
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FILE 'HOME' ENTERED AT 17:22:43 ON 06 AUG 2004

=> file caplus uspatful japio europatful medline biosis embase scisearch		
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=> s (mucosal? or transmucosal?) and aqueous?
 L1 17170 (MUCOSAL? OR TRANSMUCOSAL?) AND AQUEOUS?

=> s ((osmotic pressure) or osmolarity)
 L2 79047 ((OSMOTIC PRESSURE) OR OSMOLARITY)

=> s l1 and l2
 L3 1205 L1 AND L2

=> s l3 and mOsm
 L4 204 L3 AND MOSM

=> s l4 and (290 mOsm)
 L5 6 L4 AND (290 MOSM)

=> d l5 1-6 ibib abs

L5 ANSWER 1 OF 6 USPATFULL on STN
 ACCESSION NUMBER: 2003:89399 USPATFULL
 TITLE: Non-peptide inhibition of T-lymphocyte activation and
 therapies related thereto
 INVENTOR(S): Chandy, K. George, Laguna Beach, CA, United States
 Wulff; Heike, Irvine, CA, United States
 Cahalan, Michael D., Laguna Beach, CA, United States
 Grismer, Stephan, Blaustein, GERMANY, FEDERAL REPUBLIC
 OF
 Rauer, Heiko J., Irvine, CA, United States
 Miller, Mark J., Brea, CA, United States
 PATENT ASSIGNEE(S): The Regents of the University of California, Oakland,
 CA, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 6541494	B1	20030401
APPLICATION INFO.:	US 2000-479391		20000106 (9)
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	GRANTED		

PRIMARY EXAMINER: Travers, Russell
LEGAL REPRESENTATIVE: Buyan, Robert D., Stout, Uxa, Buyan & Mullins, LLP
NUMBER OF CLAIMS: 11
EXEMPLARY CLAIM: 1
NUMBER OF DRAWINGS: 5 Drawing Figure(s); 5 Drawing Page(s)
LINE COUNT: 1387

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Compounds, preparations and methods for immunosuppressive treatment of autoimmune disorders, graft rejection and/or graft/host disease. Therapeutically effective amounts of certain substituted triarylmethane compounds, such as 1-[(2-chlorophenyl)diphenylmethyl]-1H-pyrazole, are administered to mammalian patients to selectively inhibit the calcium-activated K^{sup.+} channel (IKCa1) in lymphocytes, monocytes, macrophages, platelets or endothelial cells without concomitant inhibition of P450-dependent enzyme systems, resulting in reduction of antigen-, cytokine-, or mitogen-induced calcium entry through store operated calcium channels in these cells, suppression of cytokine production by these cells, and inhibition of activation of these cells. Such inhibition of the Ca^{sup.++} activated K^{sup.+} channel (IKCa1) prevents the pre-Ca^{sup.++} stage of cell activation and thus causes immunosuppression and an anti-inflammatory response.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L5 ANSWER 2 OF 6 USPATFULL on STN

ACCESSION NUMBER: 2002:295175 USPATFULL
TITLE: Mediators of hedgehog signaling pathways, compositions and uses related thereto
INVENTOR(S): Baxter, Anthony David, Hertfordshire, UNITED KINGDOM
Boyd, Edward Andrew, Oxfordshire, UNITED KINGDOM
Guicherit, Oivin M., Belmont, MA, UNITED STATES
Price, Stephen, Buckinghamshire, UNITED KINGDOM
Rubin, Lee L., Wellesley, MA, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002165221	A1	20021107
APPLICATION INFO.:	US 2001-977096	A1	20011012 (9)

	NUMBER	DATE
PRIORITY INFORMATION:	US 2000-240536P	20001013 (60)
	US 2000-240564P	20001013 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	ROPES & GRAY, ONE INTERNATIONAL PLACE, BOSTON, MA, 02110-2624	
NUMBER OF CLAIMS:	92	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	58 Drawing Page(s)	
LINE COUNT:	5140	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention makes available methods and reagents for inhibiting aberrant growth states resulting from hedgehog gain-of-function, ptc loss-of-function or smoothened gain-of-function comprising contacting the cell with a hedgehog antagonist, such as a small molecule, in a sufficient amount to aberrant growth state, e.g., to agonize a normal ptc pathway or antagonize smoothened or hedgehog activity.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L5 ANSWER 3 OF 6 USPATFULL on STN

ACCESSION NUMBER: 2002:105654 USPATFULL
TITLE: Nicotine mucosal spray

INVENTOR(S): Jones, Richard L., Edmonton, CANADA

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002054856	A1	20020509
	US 6596740	B2	20030722
APPLICATION INFO.:	US 2001-983554	A1	20011024 (9)

	NUMBER	DATE
PRIORITY INFORMATION:	US 2000-243205P	20001024 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	Anita Nador, Bereskin & Parr, 40 King Street West, Box 401, Toronto, ON, M5H 3Y2	
NUMBER OF CLAIMS:	20	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	5 Drawing Page(s)	
LINE COUNT:	737	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB A composition for administration to the nasal mucosa of a subject comprises a solution of nicotine or a pharmaceutically acceptable salt thereof in a pharmaceutically acceptable solvent. The composition has a nicotine concentration less than 10 mg/ml. The composition used alone assists in reduction of the desire of a subject to smoke tobacco. It also reduces the nasal symptoms associated with administration of higher concentrations of nicotine to the nasal mucosa.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L5 ANSWER 4 OF 6 USPATFULL on STN

ACCESSION NUMBER: 93:7131 USPATFULL
TITLE: Treatment of osmotic disturbance with organic osmolytes
INVENTOR(S): Gullans, Steven R., Natick, MA, United States
Heilig, Charles W., Needham, MA, United States
PATENT ASSIGNEE(S): Brigham and Women's Hospital, Boston, MA, United States (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 5182299		19930126
APPLICATION INFO.:	US 1991-670779		19910319 (7)
RELATED APPLN. INFO.:	Continuation-in-part of Ser. No. US 1990-495575, filed on 19 Mar 1990, now abandoned		
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	Granted		
PRIMARY EXAMINER:	Raymond, Richard L.		
ASSISTANT EXAMINER:	Hollinden, Gary E.		
LEGAL REPRESENTATIVE:	Sterne, Kessler, Goldstein & Fox		
NUMBER OF CLAIMS:	14		
EXEMPLARY CLAIM:	1		
NUMBER OF DRAWINGS:	26 Drawing Figure(s); 23 Drawing Page(s)		
LINE COUNT:	2185		

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB A method of treating an osmotic disturbance in an animal which comprises administering to an animal an effective amount of an organic osmolyte, wherein the organic osmolyte is a polyol. Specific polyols include myo-inositol and sorbitol. Also included are precursors of organic osmolytes including precursors of polyols. Other polyol precursors are selected from the group consisting of glucose, glucose polymers, and glycerol. Also inclu

This invention was funded by a research grant from the National Institutes of Health, 1R01 DK36031, which provides to the United States Government certain rights in the invention.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L5 ANSWER 5 OF 6 JAPIO (C) 2004 JPO on STN
ACCESSION NUMBER: 2004-099624 JAPIO
TITLE: MEDICINAL COMPOSITION FOR ADMINISTRATION TO MUCOSA
INVENTOR: NISHIBE YOSHIHISA; KINOSHITA WATARU; KAWABE HIROYUKI
PATENT ASSIGNEE(S): TEIJIN LTD
PATENT INFORMATION:

PATENT NO	KIND	DATE	ERA	MAIN IPC
JP 2004099624	A	20040402	Heisei	A61K009-10

APPLICATION INFORMATION

STN FORMAT: JP 2003-430651 20031225
ORIGINAL: JP2003430651 Heisei
PRIORITY APPLN. INFO.: JP 1998-110887 19980421
PRIORITY APPLN. INFO.: JP 1998-110888 19980421
SOURCE: PATENT ABSTRACTS OF JAPAN (CD-ROM), Unexamined
Applications, Vol. 2004

AN 2004-099624 JAPIO

AB PROBLEM TO BE SOLVED: To provide a new medicinal composition for
administration to mucosa and having low **osmotic pressure**

SOLUTION: The medicinal composition for administration to mucosa and for
application to pharmacotherapy comprises a water-insoluble and/or slightly
water-soluble material and an **aqueous** medium and has less than
290 mOsm osmotic pressure. The
composition has superior transitivity in blood to conventional medicinal
compositions for administration to musca. And a composition for
administration to mucosa, comprising a hemostatic agent and a medicament,
is provided. The composition has superior osmosis and retention in the
mucosal tissue to the conventional medicinal compositions.
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L5 ANSWER 6 OF 6 EUROPATFULL COPYRIGHT 2004 WILA on STN

GRANTED PATENT - ERTEILTES PATENT - BREVET DELIVRE

ACCESSION NUMBER: 619729 EUROPATFULL EW 199928 FS PS
TITLE: COMPOSITION TO HELP STOP SMOKING.
ZUSAMMENSETZUNG ZUR RAUCHERENTWoeHNUNG.
COMPOSITION ET PROCEDE DE SUPPLEANCE AU TABAGISME.
INVENTOR(S): JONES, Richard, L., 10928 - 81 Street, Edmonton, Alberta
T5H 1L5, CA
PATENT ASSIGNEE(S): Pharmacia & Upjohn Aktiebolag, 112 87 Stockholm, SE
PATENT ASSIGNEE NO: 1720686
AGENT: Bannerman, David Gardner et al, Withers & Rogers,
Goldings House, 2 Hays Lane, London SE1 2HW, GB
AGENT NUMBER: 28001
OTHER SOURCE: EPB1999039 EP 0619729 B1 990714
SOURCE: Wila-EPS-1999-H28-T1
DOCUMENT TYPE: Patent
LANGUAGE: Anmeldung in Englisch; Veroeffentlichung in Englisch
DESIGNATED STATES: R AT; R BE; R CH; R DE; R DK; R ES; R FR; R GB; R GR; R
IE; R IT; R LI; R LU; R MC; R NL; R PT; R SE
PATENT INFO.PUB.TYPE: EPB1 EUROPAEISCHE PATENTSCHRIFT (Internationale
Anmeldung)

PATENT INFORMATION:

	PATENT NO	KIND	DATE
	EP 619729	B1	19990714
'OFFENLEGUNGS' DATE:			19941019
APPLICATION INFO.:	EP 1993-901989		19930104

PRIORITY APPLN. INFO.: GB 1992-47 19920103
 RELATED DOC. INFO.: WO 93-CA3 930104 INTAKZ
 WO 9312764 930708 INTPNR
 REFERENCE PAT. INFO.: DE 3241437 A GB 2030862 A
 GB 2133691 A US 4920989 A
 US 4953572 A
 REF. NON-PATENT-LIT.: DATABASE WPIL Week 8414, Derwent Publications Ltd.,
 London, GB; AN 84-086173 (14)

=> d 15 3 HIT

L5 ANSWER 3 OF 6 USPATFULL on STN

TI Nicotine mucosal spray

SUMM [0002] The invention relates to the field of nicotine mucosal sprays, to compositions comprising nicotine that can be administered in a mucosal spray form and to methods and uses therefore. In one embodiment, the invention relates to compositions and methods useful for subjects who wish to reduce tobacco smoking.

DETD [0030] The term "pharmaceutically acceptable solven." means a solvent that is physiologically tolerable at the dosage administered. Nicotine is soluble in water but an aqueous solvent should have it's pH adjusted with buffers and it's osmolarity adjusted into the physiological range. Those skilled in the art will know how to accomplish these adjustments.

DETD [0035] It is desirable that nasal administration of nicotine provides a sufficient dose of nicotine to a sufficiently large area of the nasal mucosa to give the desired rapid increase in blood nicotine level without providing a local nicotine concentration so high that it causes mucosal irritation and without requiring the delivery of such a large volume of nicotine-containing composition that a portion of the administered dose runs from the nose, causing annoyance and inconvenience to the user.

DETD [0052] Five concentrations of NNS were prepared from base solution containing 10 mg/ml nicotine. The (-) isomer of nicotine (the naturally occurring form) was obtained from Sigma-Aldrich Canada Ltd (Oakville, ON, Cat# N3876). To make the 10 mg/ml nicotine solution, 1 gm of nicotine was dissolved in 100 ml phosphate-buffered saline. Phosphate-buffered saline (PBS) was prepared by adding 0.71 gm Na.sub.2HPO.sub.4 and 0.69 gm Na.sub.2HPO.sub.4 to sterile distilled H.sub.2O to make 10 ml. Then 0.92 gm NaCl was added. The resulting solution had a pH of 6.8 and a 1 osmolarity of 290 mOsm. The five lower nicotine concentrations contained 9, 7, 5, 3 and 1 mg/ml of nicotine. The 9 mg/ml solution was prepared by diluting 18 ml of the base solution (10 mg/ml) with 2 ml PBS. The 7 mg/ml solution contained 14 ml of the base solution and 6 ml of PBS, the 5 mg/ml solution contained 14 ml of the base solution and 14 ml PBS, the 3 mg/ml solution contained 6 ml of the base solution and 14 ml PBS and the 1 mg/ml solution contained 2 ml of the base and 18 ml PBS. The viscosity of the solutions was near 1.0 cps (similar to water).

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FILE 'CAPLUS, USPATFULL, JAPIO, EUROPATFULL, MEDLINE, BIOSIS, EMBASE, SCISEARCH' ENTERED AT 17:23:12 ON 06 AUG 2004

L1 17170 S (MUCOSAL? OR TRANSMUCOSAL?) AND AQUEOUS?
 L2 79047 S ((OSMOTIC PRESSURE) OR OSMOLARITY)
 L3 1205 S L1 AND L2
 L4 204 S L3 AND MOSM
 L5 6 S L4 AND (290 MOSM)

=> s l4 and bioavailability
L6 135 L4 AND BIOAVAILABILITY

=> s l6 and hypotonic?
L7 4 L6 AND HYPOTONIC?

=> d l7 1-4 ibib abs

L7 ANSWER 1 OF 4 USPATFULL on STN

ACCESSION NUMBER: 2003:194115 USPATFULL
TITLE: Monobactam compositions and methods of use thereof
INVENTOR(S): Shawar, Ribhi M., Bellevue, WA, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2003133925	A1	20030717
APPLICATION INFO.:	US 2002-256341	A1	20020926 (10)

	NUMBER	DATE
PRIORITY INFORMATION:	US 2001-325933P	20010928 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	Chiron Corporation, Intellectual Property, P.O. Box 8097, Emeryville, CA, 94662-8097	
NUMBER OF CLAIMS:	15	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	4 Drawing Page(s)	
LINE COUNT:	849	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Methods, compounds and compositions are provided for inhibiting the growth of pathogenic microbes in vitro and of treatment of pathogenic bacterial infections in vivo using an antibacterial monobactam compound and a mucolytic agent.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 2 OF 4 USPATFULL on STN

ACCESSION NUMBER: 2003:187438 USPATFULL
TITLE: Topical formulations of natamycin/pimaricin
INVENTOR(S): Andersson, Borje S., Houston, TX, UNITED STATES
PATENT ASSIGNEE(S): Board of Regents, The University of Texas System (U.S. corporation)

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2003129225	A1	20030710
APPLICATION INFO.:	US 2002-294491	A1	20021114 (10)

	NUMBER	DATE
PRIORITY INFORMATION:	US 2001-332806P	20011114 (60)
DOCUMENT TYPE:	Utility	
FILE SEGMENT:	APPLICATION	
LEGAL REPRESENTATIVE:	FULBRIGHT & JAWORSKI L.L.P., A REGISTERED LIMITED LIABILITY PARTNERSHIP, 600 CONGRESS AVENUE, SUITE 2400, AUSTIN, TX, 78701-3271	
NUMBER OF CLAIMS:	36	
EXEMPLARY CLAIM:	1	
NUMBER OF DRAWINGS:	1 Drawing Page(s)	
LINE COUNT:	1498	

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB The present invention relates to formulations of Pimaricin (also called Natamycin), that are useful for the treatment and suppression of topical infections such as those caused by various pathogens including molds and

yeast, that are resistant to azole compounds and to Amphotericin B.
Methods for treatment of infections are also set forth.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 3 OF 4 USPATFULL on STN

ACCESSION NUMBER: 2002:192071 USPATFULL
TITLE: FORMULATED NUCLEIC ACID COMPOSITIONS AND METHODS OF
ADMINISTERING THE SAME FOR GENE THERAPY
INVENTOR(S): ROLLAND, ALAIN, THE WOODLAND, TX, UNITED STATES
MUMPER, RUSSELL J., THE WOODLAND, TX, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002103142	A1	20020801
	US 6514947	B2	20030204
APPLICATION INFO.:	US 1997-798974	A1	19970211 (8)
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	APPLICATION		
LEGAL REPRESENTATIVE:	LYON & LYON LLP/ VALENTIS INC., 633 WEST FIFTH STREET, SUITE 4700, LOS ANGELES, CA, 90071-2066		
NUMBER OF CLAIMS:	55		
EXEMPLARY CLAIM:	1		
NUMBER OF DRAWINGS:	11 Drawing Page(s)		
LINE COUNT:	2413		

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Compositions and methods for administering nucleic acid compositions in vitro to cells in culture or in vivo to an organism whereby the uptake of nucleic acids is enhanced are provided. Various compositions, including those incorporating protective, interactive, non-condensing compounds, are utilized to protect and administered nucleic acid formulation, thereby prolonging the localized bioavailability of the administered nucleic acid and enhancing expression from the nucleic acid.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L7 ANSWER 4 OF 4 USPATFULL on STN

ACCESSION NUMBER: 2002:119301 USPATFULL
TITLE: Aerosolized anti-infectives, anti-inflammatories, and
decongestants for the treatment of sinusitis
INVENTOR(S): Osbakken, Robert S., Camarillo, CA, UNITED STATES
Hale, Mary Anne, Woodland Hills, CA, UNITED STATES
Leivo, Frederick T., Carpinteria, CA, UNITED STATES
Munk, James D., Camarillo, CA, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2002061281	A1	20020523
APPLICATION INFO.:	US 2001-942959	A1	20010831 (9)
RELATED APPLN. INFO.:	Continuation-in-part of Ser. No. WO 2000-US18410, filed on 5 Jul 2000, UNKNOWN Continuation-in-part of Ser. No. US 2000-577623, filed on 25 May 2000, PENDING		

	NUMBER	DATE
PRIORITY INFORMATION:	US 1999-142618P	19990706 (60)
	US 1999-142620P	19990706 (60)
	US 1999-142621P	19990706 (60)
	US 1999-142622P	19990706 (60)
	US 1999-142624P	19990706 (60)
	US 1999-142741P	19990706 (60)
	US 1999-142881P	19990706 (60)
	US 2000-193507P	20000403 (60)
	US 2000-193508P	20000403 (60)

US 2000-193509P 20000403 (60)
US 2000-193510P 20000403 (60)
US 2000-194078P 20000403 (60)

DOCUMENT TYPE: Utility
FILE SEGMENT: APPLICATION
LEGAL REPRESENTATIVE: MORGAN, LEWIS & BOCKIUS, 1800 M STREET NW, WASHINGTON,
DC, 20036-5869

NUMBER OF CLAIMS: 37
EXEMPLARY CLAIM: 1
NUMBER OF DRAWINGS: 1 Drawing Page(s)
LINE COUNT: 1893

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Pharmaceutical compositions are described that comprise one or more active ingredients selected from the group consisting of an anti-infective agent, anti-inflammatory agent, anti-mucolytic agent, antihistamine, an antiseptic, and antibiotic combinations or combinations of others of these classes of ingredients, and particularly to compositions formulated as a solution or suspension in a unit dose for aerosol administration to treat chronic sinusitis.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

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